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(54) Title: ASSAY FOR MEASURING ACETYLATION OR DEACETYLATION ACTIVITY OF AN ENZYME

(57) Abstract: This invention is directed to a continuous method for measuring the activity of an enzyme that catalyzes the addition of an acetyl group to a residue capable of being acetylated or an enzyme that catalyzes the removal of an acetyl group from an acetylated residue. In particular the present invention is directed to a continuous method for measuring the activity of histone acetyltransferases and histone deacetylase enzymes

## ASSAY FOR MEASURING ACETYLATION OR DEACETYLATION ACTIVITY OF AN ENZYME

### Field of the Invention

5           This invention is directed to a continuous method for measuring the activity of an enzyme that catalyzes the addition of an acetyl group to a residue capable of being acetylated or an enzyme that catalyzes the removal of an acetyl group from an acetylated residue. In particular the present invention is directed to a continuous method for measuring the activity of histone acetyltransferases and histone deacetylase enzymes.

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### State of the Art

          Acetylation and deacetylation of histone proteins, transcription factors, and related proteins play a major role in the control of cellular processes. In particular, the acetylation state of histones controls how tightly the histone proteins interact with DNA, and therefore how accessible the DNA is to transcription factors. Enzymes that add acetyl groups to histones or other proteins are called histone acetyltransferases (HATs). Enzymes that remove the acetyl groups fall into two families: the histone deacetylases (HDACs) and the Sir2 family of deacetylases. Currently there are eleven known members of the mammalian HDAC family (Gray and Ekström, *Exper. Cell Res.* 2001, 262, 75-83; Zhou, et al. *Proc. Natl. Acad. Sci. USA* 2001, 98, 10572-10577; Kao et al. *J. Biol. Chem.* 2002, 277, 187-193; Gao et al. *J. Biol. Chem.* 2002, 277, 25748--25755) and seven members of the Sir2 family (Gray and Ekström, *Exper. Cell Res.* 2001, 262, 75-83).

          Histone acetyltransferases catalyze the transfer of an acetyl group from acetyl-CoA to the  $\epsilon$ -amino group of a lysine residue on the target protein. Many HAT enzymes have been characterized from eukaryotic organisms (Sterner and Berger, *Microbiol. Mol. Biol. Rev.* 2000, 64, 435-459). HDAC enzymes utilize a zinc ion at the active site of the protein to catalyze the removal of the acetyl group from acetyllysine in the form of acetate. Members of the Sir2 family of enzymes use NAD as a cofactor in the hydrolysis of acetyllysine.

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The acetylation state of histone proteins plays a major role in gene expression and in cell-cycle control, and appears to play a role in certain forms of cancer. In particular, abnormal recruitment of histone deacetylases by corepressor proteins has been shown to promote the development of promyelocytic leukemia. In tumor cell lines, several studies  
5 have shown that treatment with HDAC inhibitors can lead to growth inhibition, growth arrest, terminal differentiation, and/or apoptosis. In vivo studies have demonstrated growth inhibition of tumors and a reduction in tumor metastasis as a result of treatment with HDAC inhibitors (Krämer et al. Trends Endocrinol. Metab. 2001, 12, 294-300).

Effective study of the enzymology and inhibition of HATs, HDACs, and Sir2  
10 enzymes depends on the availability of robust assays capable of being performed in a high-throughput manner. Several assay methodologies have been developed for these enzymes, with varying degrees of utility for inhibitor screening.

Histone acetyltransferase assays are typically radioactivity-based. In these formats, acetyl-CoA radiolabeled on the acetyl group is reacted with a peptide  
15 corresponding to a histone amino acid sequence. Transfer of radiolabeled acetate to the peptide is quantitated by binding of the peptide to affinity resin (Ait-Si-Ali et al. Nucleic Acids Res. 1998, 26, 3869-3870), phosphocellulose paper (Tanner et al. J. Biol. Chem. 1999, 274, 18157-18160), or scintillation microplates (Wynne Aherne et al. Methods 2002, 26, 245-53) and measurement of the associated radioactivity. In a non-radioactive  
20 coupled assay format, the free CoA formed in the acetyltransferase reaction serves as a substrate for  $\alpha$ -ketoglutarate dehydrogenase or pyruvate dehydrogenase. Formation of NADH serves as a measure of the rate of acetyltransferase activity (Kim et al. Anal. Biochem. 2000, 280, 308-314).

The most common deacetylase assay methodology involves labeling lysine  
25 groups in histone peptides with radiolabeled acetate. The deacetylase enzyme removes the acetyl group as acetate, which is subsequently isolated by extraction and quantitated on the basis of its radioactivity (Inoue and Fujimoto, Biochim. Biophys. Acta 1970, 220, 307-316). In a variant of this approach, a scintillation proximity assay, peptides derivatized with radiolabeled acetyl groups are attached to a bead containing scintillant  
30 that emits light upon exposure to radiation. In this assay format, cleavage of the acetyl groups causes a decrease in the light emission from the scintillant (Nare, et al., Anal.

Biochem. 1999, 267, 390-396). A non-radioactivity-based assay uses peptides containing an acetyllysine group and a fluorescent tag. Reactivity is measured by high-performance liquid chromatography, using the difference in retention time of the acetylated and non-acetylated peptides to isolate and quantitate the reaction products (Hoffmann et al.

5 Nucleic Acids Res. 1999, 27, 2057-8; Hoffmann et al. Bioconjug Chem. 2001, 12, 51-5; Hoffmann et al. Arch Pharm (Weinheim) 2001, 334, 248-52). A commercial assay uses a two-step detection protocol. In the first step, a peptide containing an acetyllysine is reacted with a deacetylase for a given period of time. Following this, the reaction is quenched and the exposed lysine is reacted with a developing agent that produces a

10 fluorophore, and the amount of deacetylated lysine is quantitated using the fluorescence of the product (Biomol, Plymouth Meeting, PA, USA). More recently, a two-step, protease-coupled assay was reported, in which a peptide was designed containing a fluorescence resonance energy transfer (FRET) donor-quencher pair and an acetyllysine. After the deacetylase reaction has been allowed to run, the reaction is quenched and the

15 amount of of deacetylated peptide is quantitated by reaction of the deacetylated peptide with a protease enzyme that cleaves specifically after lysine residues (Frey et al. Presented at 224th National Meeting of the American Chemical Society, Boston, MA, August 2002; paper MEDI-121). To date, no continuous, non-radioactive histone deacetylase assays have been reported.

20 Features of the above assay formats limit their utility. Assays based on radioactivity tend to be costly, and require special handling precautions. Also, they are often difficult to perform in a high-throughput manner. Assays that measure activity on the basis of the disappearance of a signal with time rather than the appearance of signal usually yield poor signal/noise and signal/background ratios. Further, with the exception

25 of the scintillation proximity assay described above, all deacetylase assays reported to date are endpoint assays, where the extent of the reaction is determined at a single time point only. The endpoint format assumes a linear rate of product formation with time. However, this assumption is not valid if the enzyme follows burst-phase or lag-phase kinetics, if the enzyme activity decreases over the course of the reaction, if the substrate

30 is significantly depleted over the course of the reaction, or if substrate or protein aggregation occurs over the course of the reaction. A continuous format, in which the

progress of the reaction is monitored as the reaction proceeds, is more robust than an endpoint format.

Coupled assays are common in the practice of enzymology. The technique has been reviewed in the literature (Rudolph, et al. *Methods Enz.* 1979, 63, 22-42). In this type of assay, the formation of the product of an enzymatic reaction is not measured directly. Rather, the product reacts further with another enzyme or chemical to form a second product that has is readily detectable using spectroscopic or other detection methodologies. Representative examples of this methodology are a horseradish peroxidase-coupled assay for L-amino acid oxidase (Ueda et al. *Toxicon* 1988, 26, 695-706), a chymotrypsin-coupled assay for prolyl isomerase (Fischer et al. *Biomed. Biochim. Acta* 1984, 43, 1101-11), and an assay for methionine aminopeptidase in which the cleavage of a thioester bond in the enzyme substrate yields a free thiol that can react with 5,5'-dithio-bis(2-nitrobenzoic acid) (DTNB) to yield a detectable chromophore (Zhou et al. *Anal. Biochem.* 2000, 280, 159-165). In a coupled enzyme system, if the second reaction is slower than the first, an endpoint format is necessary. The first reaction must be quenched, and the second reaction is used to quantify the amount of product formed. In the case where the second reaction is more rapid, both reactions can be run at the same time. A steady-state can be achieved in which the rate of production of the first product will equal the rate of its conversion into the second product. Under this condition, the signal produced by the formation of the second product will be a measure of the formation of the first product, and thus a measure of the activity of the first enzyme. This allows for a continuous assay format, in which the enzymatic activity can be monitored directly as a function of time.

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#### SUMMARY OF THE INVENTION

In one aspect, this invention is directed to a continuous method for measuring the activity of an enzyme that catalyzes (a) the addition of an acetyl group to a residue capable of being acetylated or (b) removal of an acetyl group from an acetylated residue which method comprises incubating said enzyme with:

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- (i) a protease;
- (ii) a polypeptide comprising:

- (a) a recognition site for the protease;
  - (b) a residue, in which the acetylation state of the residue modifies the rate of cleavage of the polypeptide by the protease; and
  - (c) at least one chemical moiety, attached to the polypeptide, that possesses an optical signal that changes upon cleavage of the polypeptide; and
- 5
- (iii) measuring the change in the optical signal.

In a second aspect, this invention is directed to a continuous method for measuring the activity of an enzyme that catalyzes the addition of an acetyl group to a lysine residue or an enzyme that catalyzes the removal of acetyl group from N<sup>ε</sup>-acetylated lysine residue which method comprises incubating said enzyme with:

10

- (i) a protease;
  - (ii) a polypeptide comprising:
    - (a) a recognition site for the protease;
    - (b) a lysine or acetyllysine residue, in which the acetylation state of the residue modifies the rate of cleavage of the polypeptide by the protease; and
    - (c) at least one chemical moiety, attached to the polypeptide, that possesses an optical signal that changes upon cleavage of the polypeptide; and
  - (iii) measuring the change in the optical signal.
- 15
- 20

Preferably, the method measures the activity of an enzyme that catalyzes the removal of acetyl group from N<sup>ε</sup>-acetylated lysine. Preferably, the enzyme is HDAC1 (SEQ. ID. NO: 1), HDAC2 (SEQ. ID. NO: 2), HDAC3 (SEQ. ID. NO: 3), HDAC4 (SEQ. ID. NO: 4), HDAC5 (SEQ. ID. NO: 5), HDAC6 (SEQ. ID. NO: 6), HDAC7 (SEQ. ID. NO: 7), HDAC8 (SEQ. ID. NO: 8), HDAC9 (SEQ. ID. NO: 9), HDAC10 (SEQ. ID. NO: 10), or HDAC11 (SEQ. ID. NO: 11), any protein with 95% or greater sequence similarity any of the said enzymes, or any fragment of any of the enzymes that retains catalytic deacetylase activity. Preferably, the enzyme is SIRT1 (SEQ. ID. NO: 12), SIRT2 (SEQ. ID. NO: 13), SIRT3 (SEQ. ID. NO: 14), SIRT4 (SEQ. ID. NO: 15), SIRT5 (SEQ. ID. NO: 16), SIRT6 (SEQ. ID. NO: 17), or SIRT7 (SEQ. ID. NO: 18), any protein with 95%

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or greater sequence similarity to any of the said enzymes, or any fragment of any of the enzymes that retains catalytic deacetylase activity. Most preferably, the enzyme is HDAC1.

5 Preferably, the protease is a member of the trypsin family of proteases. More preferably, the protease is trypsin or thrombin.

Preferably, the optical signal arises from a fluorescent moiety, or optical absorption, or from a fluorescence resonance energy transfer.

Preferably, polypeptide is less than or equal to 8 or 20 amino acids in length. More preferably, the polypeptide is acetyl-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-AMC or (2-aminobenzoyl)-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-Ala-Ala-(3-(2,4-dinitrophenyl)-2,3-  
10 diaminopropionamide). Even more preferably acetyl-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-AMC.

In a third aspect, this invention is directed to a continuous method for measuring the inhibitory properties of a test compound towards the activity of an enzyme that catalyzes (a) the addition of an acetyl group to a residue capable of being acetylated or  
15 (b) removal of an acetyl group from an acetylated residue which method comprises incubating said enzyme with:

- (i) a protease;
- (ii) a polypeptide comprising:
  - (a) a recognition site for the protease;
  - (b) a residue, in which the acetylation state of the residue  
20 modifies the rate of cleavage of the polypeptide by the protease; and
  - (c) at least one chemical moiety, attached to the polypeptide, that possesses an optical signal that changes upon cleavage of the polypeptide;

25 in the presence of the test compound;

(iii) measuring the rate of increase in the optical signal wherein the difference between the rate of increase of the optical signal in the presence of the test compound and the rate of increase of the optical signal in the absence of the test compound is indicative of inhibitory properties of the test compound.

30 Preferably, this invention is directed to a continuous method for measuring the inhibitory properties of a test compound towards an enzyme that catalyzes the

addition or removal of acetyl group from N<sup>ε</sup>-acetylated lysine residue which method comprises incubating said enzyme with:

- (i) a protease;
- (ii) a polypeptide comprising:
  - 5 (a) a recognition site for the protease;
  - (b) a lysine or acetyllysine residue, in which the acetylation state of the lysine residue modifies the rate of cleavage of the polypeptide by the protease; and
  - (c) at least one chemical moiety, attached to the polypeptide,
  - 10 that possesses an optical signal that changes upon cleavage of the polypeptide;

in the presence and absence of the test compound;

- (iii) measuring the rate of increase in the optical signals wherein the difference in the rate of increase of the optical signal in the presence and absence of the test
- 15 compound is indicative of inhibitory properties of the test compound.

More preferably, the invention is directed to a continuous method for measuring the inhibitory properties of a test compound towards a histone deacetylase enzyme which method comprises incubating the histone deacetylase enzyme with:

- (i) trypsin;
- 20 (ii) acetyl-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-AMC;

in the presence and absence of the test compound; and

- (iii) measuring the increase in fluorescence at 460 nm over time, using an excitation wavelength of 355 nm wherein the difference in the rate of increase of the fluorescence in the presence and absence of the test compound is indicative of inhibitory
- 25 properties of the test compound.

Even more preferably, the histone deacetylase enzyme is HDAC1.

Particularly preferably, HDAC1 is incubated with the test compound for at least 5 minutes prior to addition of trypsin and acetyl-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-AMC.

## 30 DEFINITIONS

Unless otherwise stated, the following terms used in the specification and claims

are defined for the purposes of this Application and have the following meanings. Other terms used in the specification and claims have meanings recognized in the art.

The term "a" as used herein means at least one.

A "polypeptide" or "peptide" as used herein is a sequence of amino acids joined  
5 through amide bonds. The amino acids may be naturally occurring or non-natural. It is known in the art that side chains of several naturally occurring amino acids may be modified by the addition of chemical functionalities comprising methyl, acetyl, or phosphate groups.

A "moiety" as used herein is a molecule or portion of a molecule that possesses an  
10 optical signal or imparts an optical property to the molecule to which it is bound. For example, the amide form of 7-amino-4-methylcoumarin (AMC) when bound to a polypeptide has a weak fluorescence with an emission maximum at 395 nm. When the amide bond is cleaved, the free 7-amino-4-methylcoumarin has a very high fluorescence intensity with an emission maximum at 460 nm. Similarly, the p-nitroaniline moiety in  
15 its amide form is colorless, but when cleaved from the polypeptide it acquires an intense yellow color.

A "continuous" assay or method as used herein is one in which the process can be monitored on a constant basis without changing the process as a result of the measurement. This includes any technique in which the sample is monitored multiple  
20 times over the course of the reaction, but the preferred definition is one in which formation of a detectable product is directly related to the activity of the enzyme and the product can be quantitated in situ without any additional liquid handling or chemical reaction steps. For example, in the HDAC assay involving acetyl-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-AMC, the rate of production of free AMC is a measure of the rate of the  
25 deacetylase reaction. The production of free AMC can be monitored constantly or at arbitrarily small time intervals on the basis of its fluorescence, and thus the assay method is continuous. The reaction does not have to be stopped to detect a signal, and there are no extraction or purification steps necessary to isolate and quantitate the products.

A "proteolytic enzyme" or "protease" as used herein is an enzyme that catalyzes  
30 the cleavage of amide bonds within a polypeptide. Trypsin and thrombin are examples of proteases.

An "optical signal" as used herein is any response to illumination of a moiety that can be used to detect or quantitate the given moiety. Absorption and fluorescence are examples of optical signals.

5 A "residue" as used herein in the context of a polypeptide is an amino acid side chain that occurs within the polypeptide chain. The residue can be naturally or non-naturally occurring. For example, hydroxymethyl, thiomethyl, are naturally occurring residues.

A "recognition site" as used herein is a sequence of amino acids within a polypeptide that allows a protease enzyme to bind to and cleave the said polypeptide.

10 Unless otherwise stated, "acetyllysine" as used herein refers to lysine acetylated at the  $\epsilon$ -amino nitrogen.

"Inhibition" as used herein is a decrease in the rate of an enzyme-catalyzed reaction as a result of a compound binding to the enzyme and disrupting the interaction of the enzyme with its substrate. For the assays described herein, inhibition of a deacetylase  
15 enzyme will result in a decrease in the magnitude of the optical signal as compared with a reaction in the absence of an inhibitor.

## EXAMPLES

### Example 1

#### 20 Synthesis of acetyl-Gly-Ala-(N-acetyl-Lys)-AMC

*tert*-Boc (N-Acetyl-Lys)-AMC (445 mg, 1 mmol, purchased from Bachem) was dissolved in 4 M HCL in dioxane to provide H-(N-acetyl-Lys)-AMC as a white solid. To a solution of H-(N-acetyl-Lys)-AMC in DMF (5 ml) was added Ac-Gly-Ala-OH (188 mg, 1 mmol) using PyBOP (520 mg, 1 mmol), HOBt (135 mg, 1 mmol), and NMM  
25 (0.296 ml, 2 mmol). The reaction mixture was stirred for 1 h and monitored by MS/LC for the presence of H-(N-acetyl-Lys)-AMC. Additional amounts of PyBOP (260 mg, 0.5 mmol), HOBt (70 mg, 0.5 mmol), and NMM (0.146 ml, 1 mmol) was added and the stirring was continued for additional 4 h after which the product was isolated in quantative yield.

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### Example 2

### Measurement of histone deacetylase activity using a FRET substrate

HDAC8 was cloned, isolated, and purified as described in the literature (Buggy, et al. Biochem. J. 2000, 350, 199-205). The peptide 2-aminobenzoyl-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-Ala-Ala-(3-dinitrophenyl-(L)-2,3-diaminopropionamide) (peptide 2) was purchased from California Peptide Research, Inc. The measurement was performed in a reaction volume of 100 μL using a 96-well assay plate. HDAC8 (approx. 400 nM final concentration) in 50 mM HEPES, 100 mM KCl, 0.001% Tween-20, 0.01% bovine serum albumin, 5% DMSO, pH 7.4, was mixed with bovine trypsin (Sigma, 50 nM final concentration) and peptide 2 (20 μM final concentration). The reaction was monitored for 1 hour in a fluorescence plate reader, using an excitation wavelength of 320 nm and a detection wavelength of 405 nm. An increase of fluorescence with time was used as the measure of reaction rate.

### Example 3

#### Determination of the inhibitory properties of chemical compounds

Measurements were performed in a reaction volume of 100 μL using 96-well assay plates. HDAC-1 (200 pM final concentration) in reaction buffer (50 mM HEPES, 100 mM KCl, 0.001% Tween-20, 5% DMSO, pH 7.4) was mixed with inhibitor at various concentrations and allowed to incubate for 30 minutes, after which trypsin and acetyl-Gly-Ala-(N-acetyl-Lys)-AMC were added to final concentrations of 50 nM and 25 μM, respectively, to initiate the reaction. Negative control reactions were performed in the absence of inhibitor in replicates of eight.

The reactions were monitored in a fluorescence plate reader. After a 30 minute lag time, the fluorescence was measured over a 30 minute time frame using an excitation wavelength of 355 nm and a detection wavelength of 460 nm. The increase in fluorescence with time was used as the measure of the reaction rate. Inhibition constants were obtained using the program BatchKi (Kuzmic et al. *Anal. Biochem.* 2000, 286, 45-50).

#### SEQUENCE IDENTIFICATION

Sequence IDs utilize the accession codes of the NCBI Reference Sequence project at the National Institutes of Health. Supporting literature references are included.

- SEQ ID. NO: 1 (Taunton, et al. Science 1996, 272, 408-411; RefSeq NP\_004955)  
SEQ ID. NO: 2 (Yang, et al. Proc. Natl. Acad. Sci. U.S.A. 1996, 93, 12845-12850;  
5 RefSeq NP\_001518)  
SEQ ID. NO: 3 (Yang et al. J. Biol. Chem. 1997, 272, 28001-28007; RefSeq  
NP\_003874)  
SEQ ID. NO: 4 (Grozinger, et al. Proc. Natl. Acad. Sci. U.S.A. 1999, 96, 4868-4873;  
RefSeq NP\_006028 )  
10 SEQ ID. NO: 5 (Grozinger, et al. Proc. Natl. Acad. Sci. U.S.A. 1999, 96, 4868-4873;  
RefSeq NP\_631944)  
SEQ ID NO: 6 (Grozinger, et al. Proc. Natl. Acad. Sci. U.S.A. 1999, 96, 4868-4873;  
RefSeq NP\_006035)  
SEQ ID. NO: 7 (Kao, et al. Genes Dev. 2000, 14, 55-66; RefSeq NP\_056216)  
15 SEQ ID. NO: 8 (Buggy, et al. Biochem. J. 2000, 350, 199-205; RefSeq NP\_060956)  
SEQ ID. NO: 9 (Zhou, et al. Proc. Natl. Acad. Sci. U.S.A. 2001, 98, 10572-10577;  
RefSeq NP\_478056)  
SEQ ID. NO: 10 (Kao, et al. J. Biol. Chem. 2002, 277, 187-193; RefSeq NP\_114408)  
SEQ ID. NO: 11 (Gao, et al. J. Biol. Chem. 2002, 277, 25748-25755; RefSeq  
20 NP\_079103)  
SEQ ID. NO: 12 (Frye, Biochem. Biophys. Res. Commun. 1999, 260, 273-279; RefSeq  
NP\_036370)  
SEQ ID. NO: 13 (Frye, Biochem. Biophys. Res. Commun. 1999, 260, 273-279; RefSeq  
NP\_036369)  
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NP\_036371)  
SEQ ID. NO: 15 (Frye, Biochem. Biophys. Res. Commun. 1999, 260, 273-279; RefSeq  
NP\_036372)  
SEQ ID. NO: 16 (Frye, Biochem. Biophys. Res. Commun. 1999, 260, 273-279; RefSeq  
30 NP\_036373)

SEQ ID. NO: 17 (Frye, Biochem. Biophys. Res. Commun. 2000, 273, 793-798; RefSeq NP\_057623)

SEQ ID NO: 18 (Frye, Biochem. Biophys. Res. Commun. 2000, 273, 793-798; RefSeq NP\_057622)

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The foregoing invention has been described in some detail by way of illustration and example, for purposes of clarity and understanding. It will be obvious to one of skill in the art that changes and modifications may be practiced within the scope of the appended claims. Therefore, it is to be understood that the above description is intended  
10 to be illustrative and not restrictive. The scope of the invention should, therefore, be determined not with reference to the above description, but should instead be determined with reference to the following appended claims, along with the full scope of equivalents to which such claims are entitled.

All patents, patent applications and publications cited in this application are  
15 hereby incorporated by reference in their entirety for all purposes to the same extent as if each individual patent, patent application or publication were so individually denoted.

What is Claimed is:

1. A continuous method for measuring the activity of an enzyme that catalyzes (a) the addition of an acetyl group to a residue capable of being acetylated or (b) removal of an acetyl group from an acetylated residue which method comprises incubating said  
5 enzyme with:
  - (i) a protease;
  - (ii) a polypeptide comprising:
    - (a) a recognition site for the protease;
    - (b) a residue, in which the acetylation state of the residue  
10 modifies the rate of cleavage of the polypeptide by the protease; and
    - (c) at least one chemical moiety, attached to the polypeptide, that possesses an optical signal that changes upon cleavage of the polypeptide; and
  - (iii) measuring the change in the optical signal.  
15
2. A continuous method for measuring the activity of an enzyme that catalyzes the addition of an acetyl group to a lysine residue or an enzyme that catalyzes the removal of  
20 acetyl group from N<sup>ε</sup>-acetylated lysine residue which method comprises incubating said enzyme with:
  - (i) a protease;
  - (ii) a polypeptide comprising:
    - (a) a recognition site for the protease;
    - (b) a lysine or acetyllysine residue, in which the acetylation  
25 state of the residue modifies the rate of cleavage of the polypeptide by the protease; and
    - (c) at least one chemical moiety, attached to the polypeptide, that possesses an optical signal that changes upon cleavage of the polypeptide; and
  - (iii) measuring the change in the optical signal.  
30

3. A continuous method for measuring the activity of an enzyme that catalyzes the removal of acetyl group from N<sup>e</sup>-acetylated lysine residue which method comprises incubating said enzyme with:
- (i) a protease;
  - 5 (ii) a polypeptide comprising:
    - (a) a recognition site for the protease;
    - (b) an acetyllysine residue, which residue when acetylated attenuates the rate of cleavage of the polypeptide by the protease; and
    - (c) at least one chemical moiety, attached to the polypeptide,
    - 10 that possesses an optical signal that changes upon cleavage of the polypeptide; and
    - (iii) measuring the change in the optical signal.
4. The method of claim 3 where the enzyme is HDAC1 (SEQ. ID. NO: 1), HDAC2 (SEQ. ID. NO: 2), HDAC3 (SEQ. ID. NO: 3), HDAC4 (SEQ. ID. NO: 4), HDAC5 (SEQ. ID. NO: 5), HDAC6 (SEQ. ID. NO: 6), HDAC7 (SEQ. ID. NO: 7), HDAC8 (SEQ. ID. NO: 8), HDAC9 (SEQ. ID. NO: 9), HDAC10 (SEQ. ID. NO: 10), or HDAC11 (SEQ. ID. NO: 11), any protein with 95% or greater sequence similarity any of the said enzymes, or any fragment of any of the enzymes that retains catalytic deacetylase activity.
- 20 5. The method of claim 3 where the enzyme is SIRT1 (SEQ. ID. NO: 12), SIRT2 (SEQ. ID. NO: 13), SIRT3 (SEQ. ID. NO: 14), SIRT4 (SEQ. ID. NO: 15), SIRT5 (SEQ. ID. NO: 16), SIRT6 (SEQ. ID. NO: 17), or SIRT7 (SEQ. ID. NO: 18), any protein with 95% or greater sequence similarity to any of the said enzymes, or any fragment of any of
- 25 the enzymes that retains catalytic deacetylase activity.
6. The method of claim 3 where the protease is a member of the trypsin family of proteases.
- 30 7. The method of claim 3 where the protease is trypsin or thrombin.

8. The method of claim 3 where the optical signal arises from a fluorescent moiety.
9. The method of claim 3 where the optical signal arises from optical absorption.
- 5 10. The method of claim 3 where the optical signal arises from a fluorescence resonance energy transfer.
11. The method of claim 7 where the polypeptide is acetyl-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-AMC.
- 10 12. The method of claim 10 where the polypeptide is (2-aminobenzoyl)-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-Ala-Ala-(3-(2,4-dinitrophenyl)-2,3-diaminopropionamide).
13. The method of claim 3 where the polypeptide is less than or equal to 20 amino  
15 acids in length.
14. The method of claim 3 where the polypeptide is less than or equal to 8 amino acids in length.
- 20 15. A continuous method for measuring the activity of a histone deacetylase enzyme which method comprises incubating the histone deacetylase enzyme with:
- (i) trypsin;
  - (ii) acetyl-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-AMC; and
  - (iii) measuring the increase in fluorescence at 460 nm over time, using an  
25 excitation wavelength of 355 nm.
16. The method of claim 15 where the histone deacetylase enzyme is HDAC1.
17. A continuous method for measuring the inhibitory properties of a test compound  
30 towards an enzyme that catalyzes the addition or removal of acetyl group from N<sup>ε</sup>-acetylated lysine residue which method comprises incubating said enzyme with:

- (i) a protease;
  - (ii) a polypeptide comprising:
    - (a) a recognition site for the protease;
    - (b) a lysine or acetyllysine residue, in which the acetylation state of the lysine residue modifies the rate of cleavage of the polypeptide by the protease; and
    - (c) at least one chemical moiety, attached to the polypeptide, that possesses an optical signal that changes upon cleavage of the polypeptide;
- 10 in the presence of the test compound;
- (iii) measuring the rate of increase in the optical signal wherein the difference between the rate of increase of the optical signal in the presence of the test compound and the rate of increase of the optical signal in the absence of the test compound is indicative of inhibitory properties of the test compound.
- 15
18. A continuous method for measuring the inhibitory properties of a test compound towards a histone deacetylase enzyme which method comprises incubating the histone deacetylase enzyme with:
- (i) trypsin;
  - (iii) acetyl-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-AMC;
- 20 in the presence and absence of the test compound; and
- (iii) measuring the increase in fluorescence at 460 nm over time, using an excitation wavelength of 355 nm wherein the difference in the rate of increase of the fluorescence in the presence and absence of the test compound is indicative of inhibitory
- 25 properties of the test compound.
19. The method of claim 18 where the histone deacetylase enzyme is HDAC1.
20. The method of claim 19 wherein HDAC1 is incubated with the test compound for
- 30 at least 5 minutes prior to addition of trypsin and acetyl-Gly-Ala-(N<sup>ε</sup>-acetyllysine)-AMC.



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(54) Title: ASSAY FOR ACETYLTTRANSFERASE OR DEACETYLASE ACTIVITY

(57) Abstract: This invention is directed to a continuous method for measuring the activity of an enzyme that catalyzes the addition of an acetyl group to a residue capable of being acetylated or an enzyme that catalyzes the removal of an acetyl group from an acetylated residue e.g. histone acetyltransferases and histone deacetylase enzymes. Said enzyme is incubated with 1) a protease, 2) a polypeptide comprising a residue in which the acetylation state of the residue modifies the rate of cleavage of the polypeptide by the protease and a moiety that possesses an optical signal that changes upon cleavage of the polypeptide.

## INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 03/03764

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C12Q1/48 C12Q1/34 C12Q1/37		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols) IPC 7 C12Q		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used) EPO-Internal, WPI Data, PAJ, CHEM ABS Data, BIOSIS, EMBASE		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 01 40506 A (MIYAZAKI TOSHIAKI ;CYCLEX CO LTD (JP); TAMAI KATSUYUKI (JP); TATSU) 7 June 2001 (2001-06-07) abstract	1-20
X,P	& EP 1 243 658 A (CYCLEX CO LTD) 25 September 2002 (2002-09-25) cited for english translation of WO-document paragraphs '0021!-'0039!,'0042!; claims --- -/--	1-20
<input checked="" type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex.		
* Special categories of cited documents : *A* document defining the general state of the art which is not considered to be of particular relevance *E* earlier document but published on or after the international filing date *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) *O* document referring to an oral disclosure, use, exhibition or other means *P* document published prior to the international filing date but later than the priority date claimed *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art *&* document member of the same patent family		
Date of the actual completion of the international search  28 August 2003		Date of mailing of the international search report  11/09/2003
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel (+31-70) 340-2040, Tx. 31 651 epo nl, Fax (+31-70) 340-3016		Authorized officer  Vadot-Van Geldre, E

## INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 03/03764

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>DATABASE BIOSIS 'Online! BIOSCIENCES INFORMATION SERVICE, PHILADELPHIA, PA, US; 1980 MATOBA T ET AL: "DIGESTIBILITY OF ACETYLATED AND SUCCINYLATED PROTEINS BY PEPSIN PANCREATIN AND SOME INTRA CELLULAR PEPTIDASES" Database accession no. PREV198171059226 XP002242280 abstract &amp; AGRICULTURAL AND BIOLOGICAL CHEMISTRY, vol. 44, no. 10, 1980, pages 2323-2328, ISSN: 0002-1369</p> <p>----</p>	1-20
A	<p>DATABASE BIOSIS 'Online! BIOSCIENCES INFORMATION SERVICE, PHILADELPHIA, PA, US; May 1998 (1998-05) WITT M W ET AL: "Influence of dietary acetylated peptides on fermentation and peptidase activities in the sheep rumen." Database accession no. PREV199800344471 XP002242279 abstract &amp; JOURNAL OF APPLIED MICROBIOLOGY, vol. 84, no. 5, May 1998 (1998-05), pages 847-851, ISSN: 1364-5072</p> <p>-----</p>	1-20

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US 03/03764

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:  
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☒ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

### Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: claims 1-10 (partially), 11, 13-14 (partially), 15-16, 17 (partially), 18-20

continuous method for measuring acetylation or deacetylation activity whereby acetyl-Gly-ala-(Ne-acetyllysine)-AMC is used.

- 1.1. Claims: claims 1-10 (partially), 12, 13-14 (partially) and 17 (partially)

continuous method for measuring acetylation or deacetylation activity whereby (2-aminobenzoyl)-Gly-ala-(Ne-acetyllysine)-Ala-Ala-(3-(2,4,-dinitrophenyl)-2,3-diaminopropionamide is used.

Please note that all inventions mentioned under item 1, although not necessarily linked by a common inventive concept, could be searched without effort justifying an additional fee.

**INTERNATIONAL SEARCH REPORT**  
Information on patent family members

International Application No  
PCT/US 03/03764

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 0140506	A	07-06-2001	JP 2001149081 A	05-06-2001
			CA 2392711 A1	07-06-2001
			EP 1243658 A1	25-09-2002
			WO 0140506 A1	07-06-2001
			US 2003082668 A1	01-05-2003